**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

Claim 1 (currently amended) A kit for intravesicular instillation to a human patient comprising~~,~~(i) a first container ~~containing~~ comprising a unit dose of a therapeutic compound selected from the group consisting of resiniferatoxin, tinyatoxin, 20-homovanillyl-mezerein ~~or~~ and 20-homovanillyl-12-deoxyphorbol-13-phenylacetate, wherein the therapeutic compound is in the form of a ~~in a~~ solution concentrate or dry powder ~~form~~ and (ii) a second container ~~containing~~ comprising a physiologically compatible diluent capable of dissolving and maintaining in solution the therapeutic compound, the volume of ~~said~~ the diluent being sufficient for intravesicular instillation of the unit dose and providing a concentration of the therapeutic compound of from about 0.05 μM to 2.0 μM upon mixing the diluent with the therapeutic compound~~, and means for combining the diluent with the stock solution or lyophilized powder under sterile conditions~~.

Claim 2 (original) The kit of claim 1, wherein the first container contains a solution of resiniferatoxin dissolved in ethanol at a concentration of from 0.5 μM to 20 μM and the second container contains 100 ml of normal saline.

Claim 3 (original) The kit of claim 1, wherein the first container contains a lyophilized power comprising from 0.005μmole to 0.2 μmole resiniferatoxin and the second container contains 100 ml of 10% (v/v) ethanol in normal saline.

Claim 4 (canceled) ~~The kit of claim 1, further comprising means for transferring an instillation dose of the therapeutic compound to a patient.~~

Claim 5 (new) The kit of claim 1, wherein concentration of the therapeutic compound is between about 0.05 μM and 1.0 μM.

Claim 6 (new) The kit of claim 1, wherein the compound is resiniferatoxin.

Claim 7 (new) The kit of claim 1, wherein the second container contains a physiologically compatible solvent comprising an aqueous ethanol mixture having less than about 20% (v/v) ethanol and from about 0-1% (w/v) non-ionic detergent.

Claim 8 (new) The kit of claim 7, wherein the solvent further comprises physiologically compatible salts.

Claim 9 (new) The kit of claim 8 wherein the solvent comprises physiological saline and a maximum of about 10% (v/v) ethanol.

Claim 10 (new) The kit of claim 7, wherein the solvent further comprises buffer salts at a pH within the normal pH range of human urine.